AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

LISTING OF CLAIMS:

- 1. (Currently Amended) A pharmaceutical composition comprising, separately or together, an efficacious amount of (i) loteprednol or a pharmaceutically effective telerable ester thereof and (ii) at least one β_2 adrenoceptor agonist for simultaneous, sequential or separate administration by inhalation in the treatment of airway disorders in mammals.
- 2. (Currently Amended) The pharmaceutical composition according to claim 1, wherein the pharmaceutically <u>effective</u> tolerable ester of loteprednol is loteprednol etabonate.
- 3. (Currently Amended) The pharmaceutical composition according to claim 1, wherein the β_2 adrenoceptor agonist is selected from the group consisting of salbutamol, reproterol, salmeterol, formoterol and their pharmaceutically acceptable tolerable salts.
- 4. (Previously Amended) The pharmaceutical composition according to claim 1, which comprises (i) loteprednol and (ii) formoterol.

Attorney's Docket No. <u>034082-005</u> Application No. <u>10/089,449</u> Page 3

- 5. (Previously Amended) The pharmaceutical composition according to claim 1, which comprises (i) loteprednol and (ii) salmeterol.
- 6. (Previously Amended) The pharmaceutical composition according to claim 1, which comprises (i) loteprednol and (ii) reproterol.
- 7. (Currently Amended) A method for the treatment of allergies and/or airway disorders, comprising administering to a patient in need of such treatment an efficacious amount of (i) loteprednol and (ii) at least one β₂ adrenoceptor agonist, if appropriate together with customary pharmaceutically acceptable excipients or vehicles, for simultaneous, sequential or separate administration.
- 8. (Currently Amended) A process for the preparation of a pharmaceutical composition for the treatment of allergies and/or airway disorders, comprising an effective amount of the active compound loteprednol and at least one β₂ adrenoceptor agonist , wherein loteprednol and the β₂ adrenoceptor agonist or the β₂ adrenoceptor agonists are mixed individually or together, if appropriate together with customary pharmaceutically acceptable excipients or vehicles, and the mixture thus obtained is converted into suitable administration forms.